Curriculum Vitae

Personal information

Rhanda ADECHINA ADEHAN

Work experience

Mar 2024 – Present
Technical Officer – Pharmaceutical Regulation
World Health Organization (WHO AFRO)
Brazzaville, Republic of Congo

- Coordinate continental joint and emergency reviews of clinical trial applications for vaccines and medicines, ensuring timeline monitoring and follow-up with Member States.
- Support regulatory harmonization initiatives aligned with WHO standards and African Medicines Agency (AMA) objectives.
- Coordinate governance processes of AVAREF statutory bodies, contributing to institutional decision-making and accountability.
- Design and deliver multi-country capacity-building programmes for regulators and ethics committees.
- Conduct regulatory system assessments using the WHO Global Benchmarking Tool (GBT) and support Institutional Development Plans.
- Initiate the establishment of a Quality Management System (QMS) to institutionalize performance and knowledge management.

Jun 2020 – Mar 2024
Head of Internal Audit & Quality Management
Benin National Regulatory Authority (ABMed)
Cotonou, Benin

- Coordinate continental joint and emergency reviews of clinical trial applications for vaccines and medicines, ensuring timeline monitoring and follow-up with Member States.
- Support regulatory harmonization initiatives aligned with WHO standards and African Medicines Agency (AMA) objectives.
- Coordinate governance processes of AVAREF statutory bodies, contributing to institutional decision-making and accountability.
- Design and deliver multi-country capacity-building programmes for regulators and ethics committees.
- Conduct regulatory system assessments using the WHO Global Benchmarking Tool (GBT) and support Institutional Development Plans.
- Initiate the establishment of a Quality Management System (QMS) to institutionalize performance and knowledge management.

Jul 2017 – Dec 2020
Clinical Lead Monitor
Bayer Healthcare SAS
Lille, France

- Led national and international clinical development projects (€20K–€900K) in oncology, cardiology, and pediatrics.
- Managed regulatory submissions, stakeholder engagement, budgets, and audit/inspection readiness (CAPA).

Nov 2016 – Jul 2017
Senior Clinical Research Associate
Bayer Healthcare SAS
Lille, France

- Led the implementation and monitoring of clinical trial protocols across multiple investigational sites.
- Ensured data quality assurance and compliance with applicable clinical trial regulations and Good Clinical Practice (GCP).
- Provided technical supervision and mentoring to junior Clinical Research Associates, strengthening team performance and compliance.

Octobre 2014 – Oct 2016
Clinical Research Associate
PRA HS
Lille, France

- Supported the set-up, conduct, and monitoring of clinical trials on behalf of pharmaceutical sponsors, covering scientific, administrative, and regulatory aspects.
- Supervised trial implementation at investigational sites to ensure protocol adherence and regulatory compliance.
- Provided operational support to investigational sites in preparation for audits and inspections.
- Delivered technical training and guidance to healthcare professionals on study protocols, Good Clinical Practice (GCP), and pharmacovigilance requirements

Janv 2013 – Sep 2014
Clinical Research Associate
DOCS Global
Lille, France

- Supported the set-up, conduct, and monitoring of clinical trials on behalf of pharmaceutical sponsors, covering scientific, administrative, and regulatory aspects.
- Supervised trial implementation at investigational sites to ensure protocol adherence and regulatory compliance.
- Provided operational support to investigational sites in preparation for audits and inspections.
- Delivered technical training and guidance to healthcare professionals on study protocols, Good Clinical Practice (GCP), and pharmacovigilance requirements.

Fev 2012 – Dec 2012
Clinical Research Associate
Fovea CRO
Paris, France

- Supported the set-up, conduct, and monitoring of clinical trials on behalf of pharmaceutical sponsors, covering scientific, administrative, and regulatory aspects.
- Supervised trial implementation at investigational sites to ensure protocol adherence and regulatory compliance.
- Provided operational support to investigational sites in preparation for audits and inspections.
- Delivered technical training and guidance to healthcare professionals on study protocols, Good Clinical Practice (GCP), and pharmacovigilance requirements.

Education and training

Sept 2009 – Sept 2011
Master of Science (MSc) Science, Technology and Health Major: Clinical Research

ILIS – Faculty of Health Engineering and Management,
University of Lille, France; ilis.univ-lille.fr

Sept 2008 – Aug 2009
Bachelor's Degree Health Engineering and management
ILIS – Faculty of Health Engineering and Management,

Sept 2006 - Aug 2008
University of Lille, France; ilis.univ-lille.fr
Bachelor of Technology Biological Engineering Major: Medical Biology and Biotechnology
IUT– University of Lille, France - iut.univ-lille.fr

Additional information

Publications

Projects

Memberships

Other Relevant Information